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## **Prevalence and in-hospital outcome of aspiration in out-of-hospital intubated trauma patients**

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# Prevalence and in-hospital outcome of aspiration in out-of-hospital intubated trauma patients

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**Objective** The aim of this study was to evaluate the prevalence for aspiration in out-of-hospital intubated trauma patients and its impact on the in-hospital outcome.

**Patients and methods** Medical records of out-of-hospital intubated trauma patients, admitted from 2009 to 2014 to a level 1 trauma center, were retrospectively reviewed. Patients younger than 18 years, treated with a supraglottic airway device, without a thorax computed tomography at admission, and with lung contusions were excluded. Two hundred and eighty-one patients were further analyzed. The definition of aspiration was based on computer tomographic findings at admission. Variables were analyzed in multivariate logistic and Cox regression analyses.

**Results** Aspiration occurred in 90 (32%) patients. The mean Injury Severity Score was significantly higher in the aspiration group ( $40 \pm 26$  vs.  $33 \pm 22$ ,  $P = 0.032$ ), whereas the mean initial Glasgow Coma Scale (GCS) on scene was significantly lower ( $5 \pm 3$  vs.  $7 \pm 4$ ,  $P < 0.001$ ). In multivariate analysis, a lower initial GCS was a significant predictor for aspiration. Pneumonia, systemic inflammatory response syndrome, and sepsis occurred at similar frequencies. The 1-day (41 vs. 23%,  $P = 0.003$ ) and the 30-day mortality (53 vs. 34%,  $P = 0.003$ ) were significantly higher in the aspiration group. In survival analysis, a lower initial GCS, a higher

Injury Severity Score, and older age were independent predictors of 30-day mortality, but not aspiration.

**Conclusion** Aspiration was frequent in out-of-hospital intubated trauma patients and was associated with higher mortality and less favorable outcome, but was not an independent predictor of mortality and morbidity. However, a low initial GCS on scene was shown to be an independent predictor for aspiration and mortality. *European Journal of Emergency Medicine* 00:000–000 Copyright © Wolters Kluwer Health, Inc. All rights reserved.

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**Keywords:** aspiration, intubation, outcome, out-of-hospital, prevalence, trauma

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## Introduction

Early intubation on scene in trauma patients is indicated according to international trauma guidelines including PHTLS, ATLS or ETC. Especially patients with traumatic head injury or shock benefit from early intubation and controlled ventilation [1–3]. Furthermore, intubation prevents pulmonary aspiration [4,5]. However, urgent intubation per se is also associated with an increased risk of aspiration [6,7]. Trauma patients are often not fasted; blood, vomitus, or swelling may make it difficult to visualize the larynx and vocal cords. In addition, difficult airway situations (face and neck trauma; constrained mouth opening; cervical spine immobilization), low Glasgow Coma Scale (GCS), obesity, difficult environmental conditions (poor light; limited space), as well as inexperienced physicians can increase the risk of aspiration [8]. Previous studies have reported incidences between 1 and 20% among various patient groups requiring emergent airway management in either the emergency department or the ICU [6,9]. In contrast, data on pulmonary aspiration in the out-of-hospital setting are

limited and the reported incidence ranges between 34 and 50% [6,7,10–12].

Possible complications after aspiration are airway obstruction, Mendelson's syndrome, pneumonia, or acute respiratory distress syndrome (ARDS), which are associated with prolonged ventilator weaning, ICU stay, and increased mortality [12–14].

The aim of this study was to evaluate the prevalence and to investigate the impact of aspiration on in-hospital outcome for out-of-hospital intubated trauma patients in a level 1 trauma center in Switzerland.

## Patients and methods

In a retrospective study, all inpatient medical records from a level 1 trauma center in Switzerland between 2009 and 2014 were reviewed. The local ethical committee approved the study (KEK-ZH-No: 2011-0382).

In Switzerland, prehospital care in case of severely injured patients is usually provided by a medical doctor

(in our area usually an anesthesia resident) and a paramedic team. Severely injured patients are transferred by ambulance or helicopter to one of 12 trauma centers. In our trauma center, a standardized clinical approach is provided in the resuscitation area, where severely injured patients are usually admitted first. The standard approach consists of a primary survey and further treatment according to ATLS or ETC is applied by the trauma staff including at least one senior and one junior anesthetist, one senior, and one junior trauma surgeon as well as several nurses. The whole-body computed tomography (CT) scan is considered the gold standard and is performed in all major trauma patients as soon as possible to evaluate their relevant injuries and determine further treatment.

All trauma patients who underwent out-of-hospital intubation during the defined study period were included ( $n=434$ ). Exclusion criteria were age younger than 18 years, supraglottic airway devices, no chest CT scan at admission, and lung contusions because of the difficulties in distinguishing contusions and aspiration on chest CT scan. A total of 281 trauma patients fulfilled the inclusion and exclusion criteria, were further analyzed, and divided into an aspiration group and a no-aspiration group. Junior and senior radiologists on call reviewed the CT scans. Classic radiographic findings in acute aspiration included ground glass opacity; perihilar, ill-defined, alveolar consolidations; multifocal patchy infiltrates; and segmental or lobar consolidation, which were usually localized to one or both lung bases. The definition of aspiration was therefore based on computer tomographic findings at admission.

We used International Classification of Diseases, 10th-revision, German Modification [15,16] codes to identify all injuries, chronic comorbidities, and acute adverse events. The International Classification of Diseases, 10th-revision0 codes were encoded by professional medical coders. The procedures were coded according to the catalogue of the Swiss Surgery Classification System [17]. The Injury Severity Score (ISS) was calculated by certified medical staff who successfully completed the Association for the Advancement of Automotive medicine Abbreviated Injury Scale course.

### Statistical analyses

Categorical data were reported in absolute numbers and percent, and numerical data as mean  $\pm$  SD. The Pearson  $\chi^2$  or Fisher's exact test was used to compare categorical data, whereas the Mann–Whitney  $U$ -test was used to compare numerical data. Explanatory variables included patient demographics (age and sex), comorbidities, and injuries (ISS and GCS). The first dependent end point (outcome) was the presence of aspiration. The secondary outcomes were 1-day as well as 30-day overall mortality. A stepwise backward multivariable logistic regression model was used to identify risk factors for aspiration and

to compare aspiration, patient, and injury factors associated with mortality. Adjusted odds ratio with corresponding 95% confidence interval (CI) were tabulated. Statistical significance was set as a two-tailed  $P$ -value of less than 0.05. To answer the question of whether aspiration is an independent risk factor for a negative in-hospital outcome, time to mortality, time to death or complications, and time to extubation were included in survival Kaplan–Meier curves and compared with the log-rank Mantel–Cox test as well as in Cox regression analyses while controlling for aspiration, age, sex, ISS, GCS, and comorbidities. Two variables (liver failure and gastroesophageal reflux disease) were excluded from the multivariate analyses as they were always associated with one of the possible outcomes. All statistical analyses were carried out by IBM SPSS Statistics 22 (SPSS Inc., Chicago, Illinois, USA).

## Results

### Demographics, prevalence of aspiration, and risk factors for aspiration

Ninety out of 281 (32%) of our trauma patients admitted to our hospital had radiologic findings associated with pulmonary aspiration. Demographics such as age, sex, and source of admission were the same in both groups (Table 1). The mean ISS was significantly higher in the aspiration group ( $40 \pm 26$  vs.  $33 \pm 22$ ,  $P=0.032$ ), whereas the mean initial GCS on scene was significantly lower ( $5 \pm 3$  vs.  $7 \pm 4$ ,  $P<0.001$ ). In terms of comorbidities, mental disorders were significantly less prevalent in the aspiration group (23 vs. 35%,  $P=0.048$ ), whereas cerebrovascular diseases were significantly more prevalent (13 vs. 5.2%,  $P=0.018$ ); other comorbidities had similar frequencies in both groups (Table 2). A lower GCS was a significant risk factor for aspiration (odds ratio 0.84; 95% CI: 0.77–0.91;  $P<0.001$ ) while controlling for age, sex, ISS, and comorbidities.

### In-hospital outcome

Surgery related complications during the hospitalization, such as wound infection, dehiscence, and hematoma, occurred significantly less frequently in the aspiration group (3.3 vs. 10.0%,  $P=0.042$ ). All other complications, in particular, pneumonia (20 vs. 17%,  $P=0.58$ ), systemic inflammatory response syndrome (4 vs. 6%,  $P=0.54$ ), and sepsis (4 vs. 4%,  $P=1.0$ ), occurred at similar frequencies. Length of mechanical ventilation was equal in both groups ( $94 \pm 184$  vs.  $90 \pm 146$  h,  $P=0.43$ ). The overall 1-day mortality (41 vs. 23%,  $P=0.003$ ) and the 30-day mortality (53 vs. 34%,  $P=0.003$ ) were significantly higher in the aspiration group, whereas the length of in-hospital stay was significantly shorter ( $9 \pm 13$  vs.  $12 \pm 14$  days,  $P=0.004$ ).

**Table 1 Overview**

Parameters	Total	No aspiration	Aspiration	P-value
Patients	281 (100)	191 (68)	90 (32)	
Sex				
Female	81 (29)	54 (28)	27 (30)	0.77
Male	200 (71)	137 (72)	63 (70)	
Age (years)	54 ± 22	53 ± 21	56 ± 22	0.23
Admission from				
Private residence	268 (95)	185 (97)	83 (92)	0.08
Nursing home	4 (1.4)	2 (1.0)	2 (2.2)	0.60
Other hospital	7 (2.5)	3 (1.6)	4 (4.4)	0.22
Other	2 (0.7)	1 (0.5)	1 (1.1)	0.54
Comorbidities	160 (57)	116 (61)	44 (49)	0.61
Initial GCS on scene	7 ± 4	7 ± 4	5 ± 3	<b>&lt; 0.001</b>
ISS	35 ± 24	33 ± 22	40 ± 26	<b>0.032</b>
Head injuries	251 (89)	168 (88)	83 (92)	0.28
Extremities injuries	129 (46)	98 (51)	31 (34)	<b>0.008</b>
Thoracic injuries	106 (38)	75 (39)	31 (34)	0.44
Spine injuries	77 (27)	57 (30)	20 (22)	0.18
Neck injuries	44 (16)	35 (18)	9 (10)	0.073
Pelvic injuries	43 (15)	33 (17)	10 (11)	0.18
Abdominal injuries	40 (14)	30 (16)	10 (11)	0.30
Length of hospital stay (days)	11 ± 14	12 ± 14	9 ± 13	<b>0.004</b>
Length of stay in ICU (h)	125 ± 193	133 ± 193	110 ± 193	0.087
Length of mechanical ventilation (h)	91 ± 159	90 ± 146	95 ± 184	0.43
Any inpatient complications (except death)	109 (39)	83 (44)	26 (29)	<b>0.019</b>
1-Day mortality	81 (29)	44 (23)	37 (41)	<b>0.003</b>
30-Day mortality	113 (40)	65 (34)	48 (53)	<b>0.003</b>

Data are reported as frequency with *n* (%) or mean ± SD.

GCS, Glasgow Coma Scale; ISS, injury severity score.

Bold indicates significant *P* values.

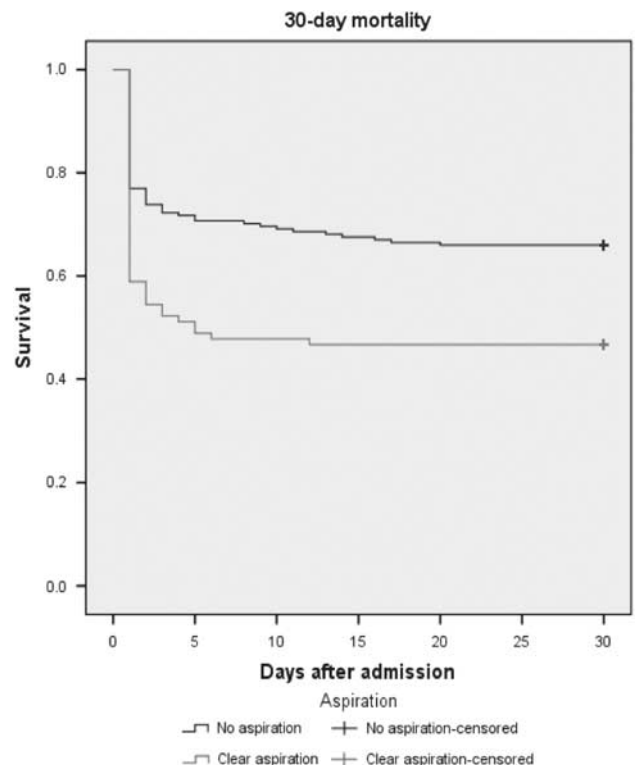
### Impact of aspiration on in-hospital outcome and other significant risk factors

In survival analyses, aspiration resulted in significantly longer time to extubation, shorter time alive and free from complications as well as higher 30-day mortality (Figs 1–3). However, in multivariate analyses, aspiration was not an independent predictor for 30-day mortality, time to death, time to extubation or time alive, and free from complications while controlling for age, sex, ISS, GCS, and comorbidities.

**Table 2 Preaccident comorbidities**

Parameters	Total [n (%)]	No aspiration [n (%)]	Aspiration [n (%)]	P-value
Mental disorders	88 (31)	67 (35)	21 (23)	<b>0.048</b>
Chronic alcoholism	24 (9)	18 (9)	6 (7)	0.44
Intoxication with psychotropic substances	22 (8)	15 (8)	7 (8)	0.98
Depression	18 (6)	12 (6)	6 (7)	0.90
Hypertensive disease	46 (16)	27 (14)	19 (21)	0.14
Arrhythmias	37 (13)	27 (14)	10 (11)	0.48
Atrial fibrillation	21 (8)	15 (8)	6 (7)	0.72
Cerebrovascular disease	22 (8)	10 (5)	12 (13)	<b>0.018</b>
Diabetes mellitus	15 (5)	12 (6)	3 (3)	0.40
Esophagitis, gastroesophageal reflux disease, peptic ulcer disease, gastritis, duodenitis	14 (5)	8 (4)	6 (7)	0.39
Muscle, skeleton and connective tissue disease	11 (4)	10 (5)	1 (1)	1.00
Paraplegia or tetraplegia and hemiplegia	8 (3)	5 (3)	3 (3)	0.71
Malignancy	7 (3)	6 (3)	1 (1)	0.44
Chronic pulmonary disease	7 (3)	6 (3)	1 (1)	0.44
Chronic coronary heart disease	6 (2)	6 (3)	0 (0)	0.18
Congestive heart failure	5 (2)	4 (2)	1 (1)	1.00

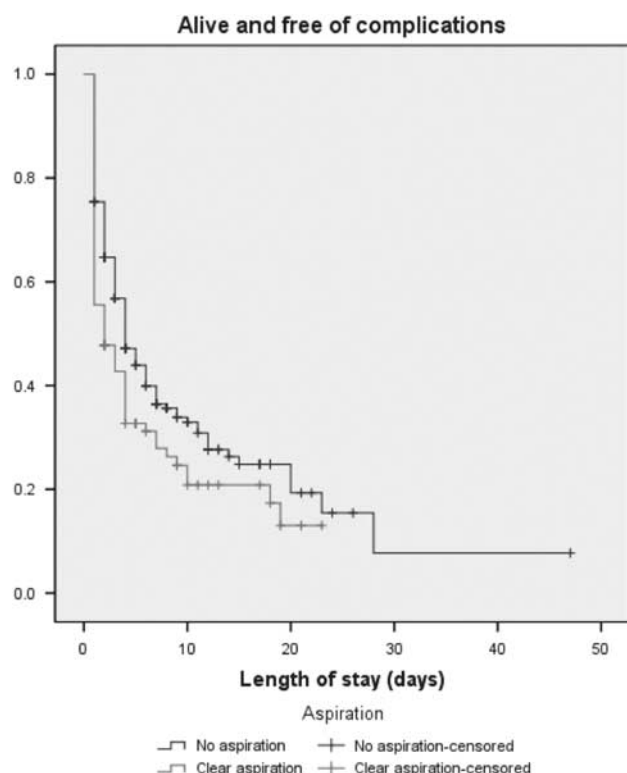
Bold indicates significant *P* values.

**Fig. 1**

30-Day mortality: mortality in trauma patients with aspiration is significantly higher compared with patients without aspiration.  $P = 0.001$  (Mantel-Cox).

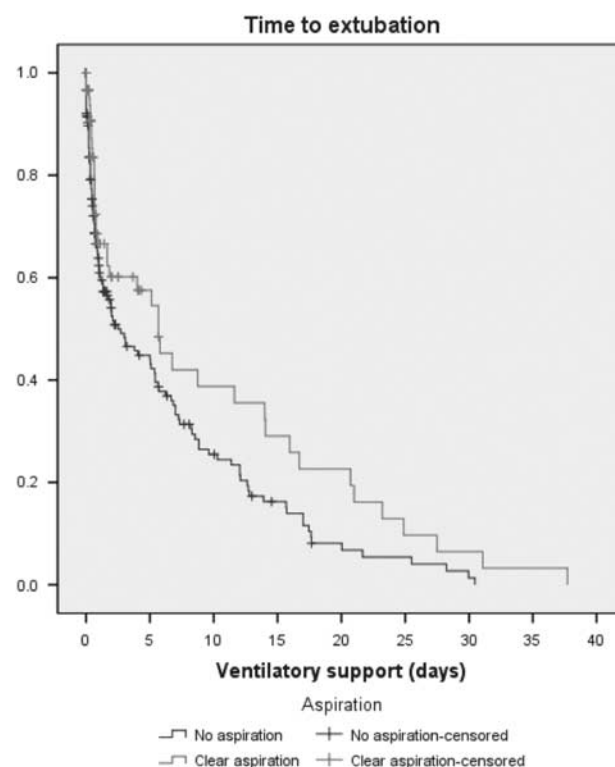
Older age, higher ISS, lower initial GCS, and inversely chronic alcoholism/mental disorders were independent predictors for 30-day mortality (Table 3) and shorter time to death. Higher ISS (HR: 1.013; 95% CI: 1.007–1.020;  $P < 0.001$ ) and older age (HR 1.008; 95% CI 1.001–1.014;  $P = 0.016$ ) were also predictors for longer time to extubation.

Fig. 2



Alive and free of complications: the time alive and free of complications is significantly shorter in patients with aspiration. Patients who died were right censored from further analysis, depicted by a vertical bar on each graph.  $P=0.028$  (Mantel–Cox).

Fig. 3



Time to extubation: trauma patients with aspiration were intubated and ventilated for a significantly longer duration. Patients who died were right censored from further analysis, depicted by a vertical bar on each graph.  $P=0.028$  (Mantel–Cox).

## Discussion

Aspiration is a common problem in the out-of-hospital airway management in trauma patients. Our study found an aspiration rate of 32% in out-of-hospital intubated trauma patients, which is slightly below the reported range from 34 to 50% [6,7,10–12]. Nevertheless, aspiration was not an independent predictor of negative in-hospital outcome.

One-third of all intubated patients showed signs of aspiration on chest CT at admission in our study. These patients with aspiration had a significantly lower initial GCS on scene, which was shown to be an independent predictor in multivariate analysis. Aspiration is commonly found to be a frequent complication of rapid sequence intubation, although particularly a longer time in a low GCS state could increase the risk of aspiration as the ability to protect the airway from aspiration decreases with the GCS. It was shown that the majority of those trauma patients who aspirated did so before the intubation because of the decreased ability to protect the airway from aspiration [12]. In addition, we carried out a sub-analysis excluding all patients with severe head injury; these results suggested that aspiration is rather the

Table 3 Independent 30-day mortality predictors

	OR	95% CI		P-value
		Lower	Upper	
ISS	1.118	1.083	1.154	<b>&lt; 0.001</b>
Age	1.033	1.014	1.052	<b>0.001</b>
Initial GCS on scene	0.74	0.64	0.85	<b>&lt; 0.001</b>
Chronic alcoholism	0.03	0.00	0.76	<b>0.033</b>

Variables entered: age, sex, ISS, GCS, aspiration, intoxication with psychotropic substances, chronic alcoholism, depression.

CI, confidence interval; GCS, Glasgow Coma Scale; ISS, injury severity score; OR, odds ratio.

Bold indicates significant  $P$  values.

consequence of a low GCS on scene than a complication because of the intubation.

Our study cohort had a high morbidity and 30-day mortality: both 40%. The high mortality rate was because of our inclusion criteria. However, higher ISS, lower GCS, and higher age were comprehensible risk factors for this adverse outcome. Mental disorders and chronic alcoholism were associated with a more favorable outcome in terms of 30-day mortality and time to death. The assessment of a proper GCS in patients with mental disorders or intoxication is challenging. In our opinion,

the initial GCS were probably lower scored in these patients by paramedics and physicians, which led to a liberal intubation strategy, although most probably indicated. The significant lower surgery-related complication rate and the shorter hospital stay in the aspiration group were rather caused by the higher mortality rate of these patients, especially in the first 24 h after admission. The higher mortality in the aspiration group might also have decreased the incidence of pneumonia in our study [18]. Pneumonia was still comparable in both groups. To the best of our knowledge, no comparable data on the impact of out-of-hospital aspiration in pneumonia have been published so far. We only found a controversial discussion on the impact on ventilator-associated pneumonia (VAP). Evans *et al.* [18] reported that out-of-hospital intubation in trauma patients was not associated with a higher risk of VAP. However, other studies have shown that rates of VAP are significantly higher (16–31%) among patients intubated in an emergency situation following trauma [12,19].

The fact that no significant increase in complications was observed in the aspiration group may also be the consequence of the current treatment strategy of these patients. Early, empiric, antibiotic therapy, controlled lung protective ventilation, and special surveillance in the intensive care unit might be a successful strategy to control typical aspiration-related complications such as pneumonia, sepsis, or ARDS. Nevertheless, aspiration is rather the consequence of a low GCS than a complication to the intubation and early, prehospital airway securing should not be under-rated as it is the first step in the prevention of aspiration and its related complications.

This study has several limitations that may have interfered with our results. First, this is a retrospective study and involves some variables that we will discuss further. Aspiration was diagnosed by the initial CT chest at admission; therefore, we have no information on the type of aspiration (blood, gastric contents, saliva, etc.) that was shown to affect pulmonary complications such as pneumonia, Mendelson syndrome, or ARDS [20]. We cannot conclude at which point the aspiration occurred: whether it was before, during, or after the intubation. In addition, we had no information on the use of drugs such as relaxants, hypnotics, and/or analgesics for rapid sequence induction and also for the use of antibiotics and their influence on in-hospital outcome. Another limitation is the out-of-hospital subjective estimation and evaluation of the patients such as GCS on scene, which led to the decision to intubation. Last but not the least, our patient population represents only traumatic injured patients from a single medical center and does not include any internal medicine patients.

## Conclusion

Aspiration was present in one of three patients; it was associated with a higher 30-day mortality and less

favorable outcome, but was not an independent predictor for mortality, time to extubation, complication, or time alive and free from complications. The current treatment strategy of trauma patients may address aspiration-related complications already in advance. However, a low initial GCS on scene was shown to be an independent predictor for aspiration and 30-day mortality.

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Authors contribution: Raluca R. Radu and Alexander Kaserer contributed to data collection, data interpretation, drafting the manuscript, and critical revision of the manuscript. Burkhardt Seifert contributed to the statistical analysis of the manuscript. Hans P. Simmen contributed to critical revision of the manuscript. Kurt Ruetzler conceived and designed the study, contributed to data interpretation, and helped to draft the manuscript. Donat R. Spahn contributed to data interpretation, drafting the manuscript, and critical revision of the manuscript. Valentin Neuhaus participated in the design and coordination of the study, carried out the statistical analysis, contributed to data interpretation, drafting the manuscript, and critical revision of the manuscript. All authors read and approved the final manuscript.

## Conflicts of interest

Dr Spahn's academic department is/has been receiving grant support from the Swiss National Science Foundation, Berne, Switzerland, the Ministry of Health (Gesundheitsdirektion) of the Canton of Zurich, Switzerland for Highly Specialized Medicine, the Swiss Society of Anesthesiology and Reanimation (SGAR), Berne, Switzerland, the Swiss Foundation for Anesthesia Research, Zurich, Switzerland, Bundesprogramm Chancengleichheit, Berne, Switzerland, CSL Behring, Berne, Switzerland, Vifor SA, Villars-sur-Glâne, Switzerland. Dr Spahn was the chair of the ABC Faculty and is the co-chair of the ABC-Trauma Faculty, managed by Physicians World Europe GmbH, Mannheim, Germany and sponsored by unrestricted educational grants from Novo Nordisk HealthCare AG, Zurich, Switzerland, CSL Behring GmbH, Marburg, Germany and LFB Biomédicaments, Courtaboeuf Cedex, France. In the past 5 years, Dr Spahn has received honoraria or travel support for consulting or lecturing from: Abbott AG, Baar, Switzerland, AMGEN GmbH, Munich, Germany, AstraZeneca AG, Zug, Switzerland, Baxter AG, Volketswil, Switzerland, Baxter S.p.A., Roma, Italy, Bayer, Zürich, Switzerland and Berlin, Germany, B. Braun Melsungen AG, Melsungen, Germany, Boehringer Ingelheim (Schweiz) GmbH, Basel, Switzerland, Bristol-Myers-Squibb, Rueil-Malmaison Cedex, France and Baar, Switzerland, CSL Behring GmbH, Hattersheim am Main, Germany and Berne, Switzerland, Curacyte AG, Munich, Germany, Daiichi Sankyo (Schweiz) AG, Thalwil, Switzerland, Ethicon Biosurgery, Sommerville, New Jersey, USA, Fresenius SE, Bad Homburg v.d.H., Germany,

Galenica AG, Bern, Switzerland (including Vifor SA, Villars-sur-Glâne, Switzerland), GlaxoSmithKline GmbH & Co. KG, Hamburg, Germany, Janssen-Cilag, Baar, Switzerland and Beers, Belgium, LFB Biomédicaments, Courtaboeuf Cedex, France, Merck Sharp & Dohme AG, Luzern, Switzerland, Novo Nordisk A/S, Bagsværd, Denmark, Octapharma AG, Lachen, Switzerland, Organon AG, Pfäffikon/SZ, Switzerland, PAION Deutschland GmbH, Aachen, Germany, Pharmacosmos A/S, Holbaek, Denmark, Photonics Healthcare B.V., Utrecht, Netherlands, ratiopharm Arzneimittel Vertriebs-GmbH, Vienna, Austria, Roche, Reinach, Switzerland, Sarstedt AG & Co., Nümbrecht, Germany, Schering-Plough International, Inc., Kenilworth, New Jersey, USA, Tem International GmbH, Munich, Germany, Verum Diagnostica GmbH, Munich, Germany, Vifor Pharma, Munich, Germany, Vienna, Austria and St. Gallen, Switzerland. For the remaining authors there are no conflicts of interest.

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